

## Guidelines for the Prevention and Treatment of Opportunistic Infections in Adults and Adolescents with HIV.

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Table 4. Indications for Discontinuing and Restarting Opportunistic Infection Secondary Prophylaxis or Chronic Maintenance in HIV-Infected Adults and Adolescents (page 1 of 3) (Last updated November 21, 2019; last reviewed November 21, 2019)

**Indication for Indication for Indication for Indication for Discontinuing Restarting Secondary Opportunistic** Discontinuina Restarting Secondary Prophylaxis/ Infection Prophylaxis/Chronic Primary Primary **Chronic Maintenance Therapy Prophylaxis Prophylaxis** Maintenance **Bacterial Enteric** Not applicable Not applicable Resolution of Salmonella infection and No recommendation Infections: after response to ART with sustained Salmonellosis viral suppression and CD4 counts >200 cells/µL (CII) **Bartonellosis** Not applicable Not applicable · Received at least 3-4 months of No recommendation treatment, and • CD4 count >200 cells/µL for ≥6 months (CIII) · Some specialists would only discontinue therapy if Bartonella titers have also decreased by fourfold (CIII). Candidiasis Not applicable Not applicable If used, reasonable to discontinue when No recommendation (Mucocutaneous) CD4 count >200 cells/µL (AIII). Coccidioidomycosis CD4 count ≥250 Restart at CD4 Only for patients with focal coccidioidal No recommendation cells/µL and with count <250 cells/ pneumonia (AII): viral suppression μL (BIII) Clinically responded to ≥ 6 months while on ART (CIII) antifungal therapy, with CD4 count ≥250 cells/mm³, and with viral suppression while on ART. Should continue monitoring for recurrence with serial chest radiographs and coccidioidal serology every 6-12 months. For patients with diffuse pulmonary (BIII), disseminated non-meningeal (BIII): Therapy is at least 12 months and usually much longer; discontinuation is dependent on clinical and serological response and should be made in consultation with experts For meningeal diseases (AII): Suppressive therapy should be continued indefinitely, even with increase in CD4 count on ART. Cryptococcal If the following criteria are fulfilled Not applicable Not applicable CD4 count <100 cells/µL Meningitis (BII): (AIII) Completed initial (induction and consolidation) therapy, and · Received at least 1 year of maintenance therapy, and · Remain asymptomatic of cryptococcal infection, and • CD4 count ≥100 cells/µL for >3 months, and with suppressed plasma HIV RNA in response to ART

Table 4. Indications for Discontinuing and Restarting Opportunistic Infection Secondary Prophylaxis or Chronic Maintenance in HIV-Infected Adults and Adolescents (page 2 of 3)

Opportunistic Infection	Indication for Discontinuing Primary Prophylaxis	Indication for Restarting Primary Prophylaxis	Indication for Discontinuing Secondary Prophylaxis/ Chronic Maintenance Therapy	Indication for Restarting Secondary Prophylaxis/Chronic Maintenance
Cystoisosporiasis (Formerly Isosporiasis)	Not applicable	Not applicable	Sustained increase in CD4 count to >200 cells/µL for >6 months in response to ART and without evidence of <i>I. belli</i> infection (BIII)	No recommendation
Cytomegalovirus Retinitis	Not applicable	Not applicable	<ul> <li>CMV treatment for at least 3 to 6 months; and with CD4 count &gt;100 cells/µL for &gt;3 to 6 months in response to ART (AII).</li> <li>Therapy should be discontinued only after consultation with an ophthalmologist, taking into account anatomic location of lesions, vision in the contralateral eye, and feasibility of regular ophthalmologic monitoring.</li> </ul>	CD4 count <100 cells/ µL (AIII)
			Routine (i.e., every 3 months) ophthalmologic follow-up is recommended after stopping therapy for early detection of relapse or immune restoration uveitis, and then periodically after sustained immune reconstitution (AIII).	
Histoplasmosis	On ART, with CD4 count >150 cells/mm³ and undetectable HIV-1 viral load for 6 months (BIII)	For patients at high risk of acquiring histoplasmosis, restart if CD4 count falls to <150 cells/ mm³ (CIII)	If the following criteria (AI) are fulfilled:  • Received azole therapy for >1 year, and  • Negative fungal blood cultures, and  • Serum or urine Histoplasma antigen below the level of quantification, and  • Undetectable HIV viral load, and  • CD4 count ≥150 cells/mm³ for ≥6 months in response to ART	CD4 count <150 cells/ mm <sup>3</sup> (BIII)
Leishmaniasis: Visceral (and possibly cutaneous leishmaniasis in immunocompro- mised patients with multiple relapses)	Not applicable	Not applicable	There is no consensus regarding when to stop secondary prophylaxis. Some investigators suggest that therapy can be stopped if CD4 count increases to >200 to 350 cells/µL for 3–6 months in response to ART, but others suggest that therapy should be continued indefinitely.	No recommendation
Microsporidiosis	Not applicable	Not applicable	No signs and symptoms of non-ocular <b>(BIII)</b> or ocular <b>(CIII)</b> microsporidiosis and CD4 count >200 cells/µL for >6 months in response to ART.	No recommendation
<i>Mycobacterium</i> avium Complex Disease	Initiation of effective ART <b>(AI)</b>	CD4 count <50 cells/mm³: only if not on fully suppressive ART (AIII)	<ul> <li>If the Following Criteria are Fulfilled (AI):</li> <li>Completed ≥12 months of therapy, and</li> <li>No signs and symptoms of MAC disease, and</li> <li>Have sustained (&gt;6 months) CD4 count &gt;100 cells/mm³ in response to ART.</li> </ul>	CD4 count <100 cells/ mm³ (AIII)

Table 4. Indications for Discontinuing and Restarting Opportunistic Infection Secondary Prophylaxis or Chronic Maintenance in HIV-Infected Adults and Adolescents (page 3 of 3)

Opportunistic Infection	Indication for Discontinuing Primary Prophylaxis	Indication for Restarting Primary Prophylaxis	Indication for Discontinuing Secondary Prophylaxis/ Chronic Maintenance Therapy	Indication for Restarting Secondary Prophylaxis/Chronic Maintenance
Pneumocystis Pneumonia	CD4 count increased from <200 to >200 cells/mm³ for >3 months in response to ART (AI)  Can consider when CD4 count is 100–200 cells/mm³ if HIV RNA remains below limits of detection for ≥3 months–6 months (BII).	CD4 count <100 cells/mm³ (AIII)  CD4 count 100–200 cells/mm³ and HIV RNA above detection limit of the assay (AIII).	CD4 count increased from <200 cells/mm³ to >200 cells/mm³ for >3 months in response to ART (BII)  Can consider when CD4 count is 100—200 cells/mm³ if HIV RNA remains below limits of detection for ≥3 months—6 months (BII).  If PCP occurs at a CD4 count >200 cells/mm³ while not on ART, discontinuation of prophylaxis can be considered once HIV RNA levels are suppressed to below limits of detection for ≥3months—6 months (CIII).  If PCP occurs at a CD4 count >200 cells/mm³ while on ART, continue PCP prophylaxis for life, regardless of how high the CD4 cell count rises as a	CD4 count <100 cells/mm³ (AIII)  CD4 count 100–200 cells/mm³ and with HIV RNA above detection limit of the assay (AIII).
Talaromycosis (Penicilliosis)	CD4 count >100 cells/mm³ for >6 months in response to ART (BII) or If achieved sustained HIV viral suppression for >6 months (BIII)	CD4 count <100 cells/mm³ (BIII)—if patient is unable to have ART, or has treatment failure without access to effective ART options, and still resides in or travels to the endemic area	consequence of ART (BIII).  CD4 count >100 cells/mm³ for ≥6 months in response to ART (BII)  or  If achieved sustained HIV viral suppression for >6 months (BIII)	CD4 count <100 cells/ mm³ (BIII)
Toxoplasma gondii Encephalitis	CD4 count increased to >200 cells/µL for >3 months in response to ART (AI)  Can consider when CD4 count 100-200 cells/µL if HIV RNA remain below limits of detection for at least 3-6 months (BII)	CD4 count <100 cells/µL, (AIII)  CD4 count 100-200 cells/µL and with HIV RNA above detection limit of the assay (AIII).	Successfully completed initial therapy, remain free of signs and symptoms of TE, and CD4 count >200 cells/µL for >6 months in response to ART (BI).	CD4 count <200 cells/ μL <b>(AIII)</b>

**Key to Acronyms:** ART = antiretroviral therapy; CD4 = CD4 T lymphocyte cell; CMV = cytomegalovirus; MAC = *Mycobacterium avium* complex; PCP = *Pneumocystis* pneumonia; TE = *Toxoplasma* encephalitis

## **Evidence Rating:**

Strength of Recommendation:

- A: Strong recommendation for the statement
- B: Moderate recommendation for the statement
- C: Optional recommendation for the statement

Quality of Evidence for the Recommendation:

- I: One or more randomized trials with clinical outcomes and/or validated laboratory endpoints
- II: One or more well-designed, nonrandomized trials or observational cohort studies with long-term clinical outcomes
- III: Expert opinion

In cases where there are no data for the prevention or treatment of an OI based on studies conducted in HIV-infected populations, but data derived from HIV-uninfected patients exist that can plausibly guide management decisions for patients with HIV/AIDS, the data will be rated as III but will be assigned recommendations of A, B, C depending on the strength of recommendation.